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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 3

Complete if Known	
Application Number	09/942,336
Filing Date	August 29, 2001
First Named Inventor	Tetsuo Ashizawa
Art Unit	1637
Examiner Name	~~Hashemi, Shar S.~~ Teresa Knelecka
Attorney Docket Number	HO-P02039US1

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number: Number-Kind Code[2] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
TS	AA	U.S. 5,853,995	12-29-1998	Lee; Cheng-Chi	
	AB	U.S. 5,840,491	11-24-1998	Kakizuka; Akira	
	AC	U.S. 5,834,183	11-10-1998	Orr; Harry T; Ranum; Laura P. W.; Chung; Ming-yi; Zoghbi; Huda Y.	
TS	AD	U.S. 5,981,185	11-09-1999	Matson et al.	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document: Country Code[3]-Number[4]-Kind Code[5] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]

[1] Applicant's unique citation designation number (optional). [2] See attached Kinds Codes of USPTO Patent Documents at www.uspto.gov or MPEP 901.04. [3] Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). [4] For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the application number of the patent document. [5] Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. [6] Applicant is to place a check mark here if English language Translation is attached.

OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
TS	CA	Timchenko, LT and Caskey, CT; Trinucleotide repeat disorders in humans: discussions of mechanisms and medical issues; FASEB J., Dec. 1996, pp. 1589-1597; vol. 10(14).	
	CB	Burgess, DL, Matasuura, T., Ashizawa, T., Noebels, JL; Genetic localization of the Ca2+ channel gene CACNG2 near SCA10 on chromosome 22q13; Epilepsia, Jan. 2000, pp. 24-27; vol. 14(1).	
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TS	CJ	Ishikawa, K., Mizusawa, H., Saito, M., Tanaka, H., et al. Autosomal dominant pure cerebellar ataxia. A clinical and genetic analysis of eight Japanese families. Brain 1996, pp. 1173-1182, vol. 119 (Pt. 4).	
	~~CK~~	~~Cedars-Sinai scientists localize new ataxia/epilepsy gene; February 17, 1999;~~	

Teresa Knelecka 7/16/03

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 3

Complete if Known	
Application Number	09/942,336
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Examiner Name	~~Hashemi, Shar S.~~ Teresa Strelecka
Attorney Docket Number	HO-P02039US1

		~~http://eurekalert.org/releases/cs-css021999.html~~	
	CL	Biros, I., Forrest, S.M. Duplex PCR for autosomal dominant spinocerebellar ataxia testing: A nonradioactive rapid screening method; Molecular Diagnosis; http://www.wbsaunders.com/MoleDiag/abs/abs3_4/00300223.html	
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	CO ✓	Holmes, S.E. et al. Expansion of a novel CAG trinucleotide repeat in the 5' region of PPP2R2B is associated with SCA12; Nature Genet. 1999, pp. 391-392, vol. 23.	
	CP ✓	Imbert, G. et al. Cloning of the gene for spinocerebellar ataxia 2 reveals a locus with high sensitivity to expanded CAG/glutamine repeats. Nature Genet. 1996, pp. 285-291, vol. 14.	
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	CR ✓	Orr H.T. et al., Expansion of an unstable trinucleotide CAG repeat in spinocerebellar ataxia type 1. Nature Genet. 1993, pp. 221-226, vol. 4.	
	CS ✓	Pulst S.M. et al., Moderate expansion of a normally biallelic trinucleotide repeat in spinocerebellar ataxia type 2. Nature Genet 1996, pp. 269-276, vol. 14.	
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TS	DA ✓	Ashizawa T, Matsuura T, Rasmussen A, Grewal RP, Zu L, Pulst SM, Pandolfo M, Sasaki H, Volpini V, Yamagata T, Watase K, Burgess DL, Inoue K, Yescas P, Nagamitsu S, Momoi MY, Tashiro K, Zoghbi HY, Alonso E, Nelson DL. Founder effect of the spinocerebellar ataxia type 10 mutation in the Mexican population. Am J Hum Genet 2000;67:373. Abstract submission faxed 7/15/03	

Examiner Signature	Teresa Strelecka	Date Considered	7/16/03

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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